

UNIVERSAL ANTI-CAVITY FLUORIDE- sodium monofluorophosphate paste
Universal Distribution Center LLC

UNIVERSAL ANTI-CAVITY FLUORIDE TOOTHPASTE

Active Ingredient

Sodium Monofluorophosphate 0.76%
(1000 ppm)

Purpose

Anticavity toothpaste

Uses

- regular brushing with flouride toothpaste helps protect teeth against cavities
- aids in the prevention of dental cavities.

Warning

When using this product

- if pain\ sensitivity still persists after 4 weeks of use, please visit your dentist.

Stop and ask a dentist

- if the problem persists or worsens.

Sensitivity teeth may indicate a serious problem that may need prompt care by a dentist.

Keep out of reach of children under 6 years of age

- If accidentally swallowed get medical help or contact a Poison Control Center right away.

Directions

- adults and children of 2 years and older: brush teeth thoroughly, after meals or at least twice a day, or as directed by a dentist.
- do not swallow.
- to minimize swallowing, use a pea-sized amount in children under 6 years old.
- supervise children's brushing until good habits are established.
- children under 2 years: ask a dentist before use.

Children under 12 years of age: Consult a dentist or doctor

Other information

- store in a cool, dry place.

Inactive Ingredients

Calcium carbonate, Water, Glycerine, Silica, Sodium Lauryl sulphate, Sorbitol, Xanthan gum, Sodium saccharine, Sodium benzoate, Sodium carboxy methyl cellulose and Flavor

PRINCIPAL DISPLAY PANEL

UNIVERSAL ANTI-CAVITY FLUORIDE TOOTHPASTE



UNIVERSAL ANTI-CAVITY FLUORIDE

sodium monofluorophosphate paste

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-111
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU4080)	FLUORIDE ION	0.76 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SORBITOL (UNII: 506T60A25R)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-111-01	181 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	05/30/2020	

Labeler - Universal Distribution Center LLC (019180459)

Revised: 11/2023

Universal Distribution Center LLC